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APPLICATION NO.	FIL	ING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/047,608	01	/14/2002	Leonard Bell	59	59 5748	
7590 12/23/2004			EXAM	EXAMINER		
Mark Farber Alexion Pharmaceuticals				VANDERVEGT, FRANCOIS P		
352 Knotter Drive				ART UNIT	PAPER NUMBER	
Cheshire, CT	06410			1644		
				DATE MAILED: 12/23/2004	DATE MAILED: 12/23/2004	

5/112 WHILEED. 12/25/200-

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)						
Advisory Action	10/047,608	BELL, LEONARD						
Advisory Action	Examiner	Art Unit						
	F. Pierre VanderVegt	1644						
The MAILING DATE of this communication appe	ars on the cover sheet with the c	orrespondence addre	ess					
THE REPLY FILED 22 November 2004 FAILS TO PLACE THIS APPLICATION IN CONDITION FOR ALLOWANCE. Therefore, further action by the applicant is required to avoid abandonment of this application. A proper reply to a final rejection under 37 CFR 1.113 may only be either: (1) a timely filed amendment which places the application in condition for allowance; (2) a timely filed Notice of Appeal (with appeal fee); or (3) a timely filed Request for Continued Examination (RCE) in compliance with 37 CFR 1.114.								
PERIOD FOR REPLY [check either a) or b)]								
a) The period for reply expiresmonths from the mailing date of the final rejection. b) The period for reply expires on: (1) the mailing date of this Advisory Action, or (2) the date set forth in the final rejection, whichever is later. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of the final rejection. ONLY CHECK THIS BOX WHEN THE FIRST REPLY WAS FILED WITHIN TWO MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f). Extensions of time may be obtained under 37 CFR 1.136(a). The date on which the petition under 37 CFR 1.136(a) and the appropriate extension fee have been filed is the date for purposes of determining the period of extension and the corresponding amount of the fee. The appropriate extension fee under 37 CFR 1.17(a) is calculated from: (1) the expiration date of the shortened statutory period for reply originally set in the final Office action; or (2) as set forth in								
(b) above, if checked. Any reply received by the Office later than three more earned patent term adjustment. See 37 CFR 1.704(b).	nths after the mailing date of the final reje	ction, even if timely filed, m	nay reduce any					
1. A Notice of Appeal was filed on <u>22 November 2004</u> . Appellant's Brief must be filed within the period set forth in 37 CFR 1.192(a), or any extension thereof (37 CFR 1.191(d)), to avoid dismissal of the appeal.								
2. The proposed amendment(s) will not be entered because:								
(a) they raise new issues that would require further consideration and/or search (see NOTE below);								
(b) they raise the issue of new matter (see Note b								
(c) they are not deemed to place the application in issues for appeal; and/or	n better form for appeal by mate	erially reducing or sir	mplifying the					
(d) they present additional claims without canceli	ing a corresponding number of f	inally rejected claims	S.					
NOTE:								
3. Applicant's reply has overcome the following reject		onarata timely filed:	amendment					
4. Newly proposed or amended claim(s) would be allowable if submitted in a separate, timely filed amendment canceling the non-allowable claim(s).								
5.⊠ The a) affidavit, b) exhibit, or c) request for reconsideration has been considered but does NOT place the application in condition for allowance because: See Continuation Sheet.								
6. The affidavit or exhibit will NOT be considered becaraised by the Examiner in the final rejection.	ause it is not directed SOLELY	to issues which were	e newly					
7. For purposes of Appeal, the proposed amendment explanation of how the new or amended claims we	(s) a) will not be entered or b) ould be rejected is provided belo	⊠ will be entered ar w or appended.	nd an					
The status of the claim(s) is (or will be) as follows:								
Claim(s) allowed:								
Claim(s) objected to:								
Claim(s) rejected: <u>1-13</u> .	Claim(s) rejected: <u>1-13</u> .							
Claim(s) withdrawn from consideration:								
8. ☐ The drawing correction filed on is a) ☐ approved or b) ☐ disapproved by the Examiner.								
9. Note the attached Information Disclosure Statemen	nt(s)(PTO-1449) Paper No(s).	 •						
10. Other:	PATRICK J. NOLAN, PRIMARY EXAMIN							
	12/22/04							

ontinuation Sheet (PTOL-303) 10/047,608

Continuation of 5. does NOT place the application in condition for allowance because: Applicant has amended the claims to recite "peak" levels of CK-MB. Applicant argues that the Fitch reference does not anticipate the claimed invention because Fitch does not relate prophylactic anti-inflammatory treatment with prevention of post-operative myocardial infarctions and because Fitch does not measure peak levels of CK-MB. However, Applicant's amendment and arguments fail to distinguish the claimed invention from the teachings of Fitch. The "peak CK-MB levels" in nanograms/ml are measurements made post-operatively, while the anti-inflammatory administration, as a prophylactic treatment, occurs pre-operatively. Both the instant specification and Fitch teach the prophylactic administration of anti-inflammatory agent. Post-operatively, Fitch measures CK-MB lelvels in I.U. and quantifies "myocardial damage." The mere fact that Fitch measures different parameters post-operatively than the instant inventors does not change the fact that the method of the instantly claimed invention and that of Fitch are the same, pre-operative administration of an anti-inflammatory compound. Measurement or observation of a different post-operative benefit does not confer patentability, but merely constitutes further characterization of an otherwise old method.